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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
10/002,784	11/26/2001	Robert G. Ulrich	003/233/SAP	6261
75	90 03/25/2003			
ATTN: MCMR-JA			EXAMINER	
U. S. Army Medical Research, and Materiel Command			MINNIFIEL	D, NITA M
504 Scott Street			ART UNIT	PAPER NUMBER
Fort Detrick, M	D 21702-5012		1645	
			DATE MAILED: 03/25/2003	10

Please find below and/or attached an Office communication concerning this application or proceeding.

	Application No.	Applicant(s)				
_	10/002,784	ULRICH, ROBERT G.				
Office Action Summary	Examiner	Art Unit				
-	N. M. Minnifield	1645				
The MAILING DATE of this communication		et with the correspondence address				
Period for Reply		4 MONTH(S) EROM				
A SHORTENED STATUTORY PERIOD FOR RE THE MAILING DATE OF THIS COMMUNICATIO - Extensions of time may be available under the provisions of 37 CFI after SIX (6) MONTHS from the mailing date of this communication - If the period for reply specified above is less than thirty (30) days, a - If NO period for reply is specified above, the maximum statutory pe - Failure to reply within the set or extended period for reply will, by st - Any reply received by the Office later than three months after the meanned patent term adjustment. See 37 CFR 1.704(b). Status	DN. R 1.136(a). In no event, however, m n. a reply within the statutory minimum priod will apply and will expire SIX (6) tetute cause the application to beco	nay a reply be timely filed of thirty (30) days will be considered timely.) MONTHS from the mailing date of this communication. me ABANDONED (35 U.S.C. § 133).				
1) Responsive to communication(s) filed on	·					
24)	This action is non-final.					
3) Since this application is in condition for al	lowance except for forma	I matters, prosecution as to the merits is				
closed in accordance with the practice un Disposition of Claims		5 G.B. 11, 400 G.G. 210.				
4)⊠ Claim(s) <u>1-75</u> is/are pending in the application.						
4a) Of the above claim(s) is/are with	ndrawn from consideration	٦.				
5) Claim(s) is/are allowed.						
6)☐ Claim(s) is/are rejected.						
7) Claim(s) is/are objected to.						
8) Claim(s) <u>1-75</u> are subject to restriction and	d/or election requirement.					
Application Papers	•					
9) The specification is objected to by the Example 1		hy the Everniner				
10) The drawing(s) filed on is/are: a) = 1						
Applicant may not request that any objection 11) The proposed drawing correction filed on _						
If approved, corrected drawings are required in reply to this Office action.						
12) The oath or declaration is objected to by the Examiner.						
Priority under 35 U.S.C. §§ 119 and 120	projen priority under 35 U	S.C. & 119(a)-(d) or (f)				
13) Acknowledgment is made of a claim for foreign priority under 35 U.S.C. § 119(a)-(d) or (f).						
a) ☐ All b) ☐ Some * c) ☐ None of:	monto hava haan receive	ч				
1. Certified copies of the priority docu						
2. Certified copies of the priority docu						
 3. Copies of the certified copies of the application from the Internation * See the attached detailed Office action for 	al Bureau (PCT Rule 17.2	2(a)).				
14) Acknowledgment is made of a claim for domestic priority under 35 U.S.C. § 119(e) (to a provisional application).						
a) ☐ The translation of the foreign languages	ge provisional application	has been received.				
Attachment(s)	•					
1) Notice of References Cited (PTO-892) 2) Notice of Draftsperson's Patent Drawing Review (PTO-94 3) Information Disclosure Statement(s) (PTO-1449) Paper N	18) 5) No	erview Summary (PTO-413) Paper No(s) bice of Informal Patent Application (PTO-152) her:				

Art Unit: 1645

DETAILED ACTION

Election/Restrictions

- 1. Restriction to one of the following inventions is required under 35 U.S.C. 121:
 - I. Claims 1-31, drawn to superantigen toxin DNA fragment and DNA construct, classified in class 536, subclass 23.1.
 - II. Claims 32-43, drawn to superantigen toxin and altered toxin, classified in class 530, subclass 350.
 - III. Claims 44 and 45, drawn to a method for diagnosis of superantigenassociated bacterial infection, classified in class 435, subclass 7.1.
 - IV. Claim 46, drawn to a diagnostic kit, classified in class 435, subclass975.
 - V. Claims 47-52, drawn to a vaccine and multivalent vaccine, classified in class 424, subclass 184.1.
 - VI. Claims 53-56, drawn to a therapeutic method for treatment or amelioration of infection, classified in class 514, subclass 2.
 - VII. Claims 57-75, drawn to antisera/antibody, classified in class 530, subclass 387.1.

The inventions are distinct, each from the other because of the following reasons:

Art Unit: 1645

The inventions of Group I, II, IV, V and VII are drawn to structurally and functionally distinct products, which are capable of separate manufacture, use and sale. Group I is directed to DNA. Group II is directed to protein. Group IV is directed to a kit. Group V is directed to a vaccine. Group VII is directed to antibody.

Inventions III and VI are unrelated. Inventions are unrelated if it can be shown that they are not disclosed as capable of use together and they have different modes of operation, different functions, or different effects (MPEP § 806.04, MPEP § 808.01). In the instant case the different inventions have different method objectives, steps and parameters.

Inventions II and III are related as product and process of use. The inventions can be shown to be distinct if either or both of the following can be shown: (1) the process for using the product as claimed can be practiced with another materially different product or (2) the product as claimed can be used in a materially different process of using that product (MPEP § 806.05(h)). In the instant case the protein can be used in a materially different process of using the product, such as treatment methods, methods to elicit antibody production or purification methods.

Art Unit: 1645

Inventions V and VI are related as product and process of use. The inventions can be shown to be distinct if either or both of the following can be shown: (1) the process for using the product as claimed can be practiced with another materially different product or (2) the product as claimed can be used in a materially different process of using that product (MPEP § 806.05(h)). In the instant case the vaccine can be used in methods to elicit specific antibody production.

Because these inventions are distinct for the reasons given above and have acquired a separate status in the art as shown by their different classification, restriction for examination purposes as indicated is proper.

Because these inventions are distinct for the reasons given above and the search required for one Group is not required for another Group, restriction for examination purposes as indicated is proper.

Because these inventions are distinct for the reasons given above and have acquired a separate status in the art because of their recognized divergent subject matter, restriction for examination purposes as indicated is proper.

Art Unit: 1645

2. This application contains claims directed to the following patentably distinct species of the claimed invention:

If Applicant elects Group I: Applicant should elect a specific superantigen toxin and SEQ ID NO.

If Applicant elects Group II: Applicant should elect a specific superantigen toxin and SEQ ID NO, as well as a specific altered superantigen toxin.

If Applicant elects Groups III-VI: Applicant should elect a specific superantigen toxin.

Applicant is required under 35 U.S.C. 121 to elect a single disclosed species for prosecution on the merits to which the claims shall be restricted if no generic claim is finally held to be allowable.

Applicant is advised that a reply to this requirement must include an identification of the species that is elected consonant with this requirement, and a listing of all claims readable thereon, including any claims subsequently added. An argument that a claim is allowable or that all claims are generic is considered nonresponsive unless accompanied by an election.

Upon the allowance of a generic claim, applicant will be entitled to consideration of claims to additional species which are written in dependent form

Application/Control Number: 10/002,784 Page 6

Art Unit: 1645

or otherwise include all the limitations of an allowed generic claim as provided by 37 CFR 1.141. If claims are added after the election, applicant must indicate which are readable upon the elected species. MPEP § 809.02(a).

Should applicant traverse on the ground that the species are not patentably distinct, applicant should submit evidence or identify such evidence now of record showing the species to be obvious variants or clearly admit on the record that this is the case. In either instance, if the examiner finds one of the inventions unpatentable over the prior art, the evidence or admission may be used in a rejection under 35 U.S.C. 103(a) of the other invention.

Applicant is advised that the reply to this requirement to be complete must include an election of the invention to be examined even though the requirement be traversed (37 CFR 1.143).

3. Any inquiry concerning this communication or earlier communications from the examiner should be directed to N. M. Minnifield whose telephone number is 703-305-3394. The examiner can normally be reached on M-F (8:00-5:30) Second Friday Off.

Application/Control Number: 10/002,784 Page 7

Art Unit: 1645

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Lynette R.F. Smith can be reached on 703-308-3909. The fax phone numbers for the organization where this application or proceeding is assigned are 703-308-4556 for regular communications and 703-308-4556 for After Final communications.

Any inquiry of a general nature or relating to the status of this application or proceeding should be directed to the receptionist whose telephone number is 703-308-0196.

Primary Examiner

APT Unit 1645

NMM March 19, 2003